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Listing of Claims:

1-12. (Canceled)

- 13. (Currently amended) A method of identifying peptoids which are effective in transfecting a cell with an oligonucleotide, the method comprising
- (i) providing a library of peptoids having a plurality of unknown different sequences and having the general formula I:

$$R^{a} \leftarrow \begin{array}{c} R^{b} & O \\ I & -CR^{1}R^{2} - C \\ I & -CR^{2} \end{array}$$

where

R^a is selected from the group consisting of alkyl, aryl, aralkyl, aralkenyl, and aralkynyl, any of which may be substituted with one or more groups X; hydrogen, -OH, -SH, -COOH, sulfonyl, and a lipid moiety, wherein said lipid moiety may be conjugated to a linker moiety,

each R^b is independently selected from the group consisting of alkyl, aryl, aralkyl, aralkenyl, and aralkynyl, any of which may be substituted with one or more groups X; and hydrogen,

wherein at least one group Rb is not hydrogen;

R° is selected from the group consisting of alkyl, aryl, aralkyl, aralkenyl, and aralkynyl, any of which may be substituted one or more groups X; hydrogen, -OH, -SH, -NH2, -NHR, -NH(C=O)R, where R is lower alkyl; sulfonyl, hydrazine, and a lipid moiety, wherein said lipid moiety may be conjugated to a linker moiety;

X is selected from hydroxy, alkoxy, amino, guanidino, amidino, alkylamino, alkylthio, halogen, nitro, cyano, keto, aldehyde, carboxylic acid, carboxylic ester, carboxylic amide, sulfonic acid and sulfonic ester;

at least one of Ra and Rc comprises a lipid moiety;

R¹ and R² are independently selected from hydrogen, lower alkyl, and lower alkoxy; and m is an integer selected from 2 to about 50, wherein

the sequences of individual peptoids in the library are unidentified;

- (ii) contacting a plurality of unknown peptoids <u>having unidentified sequences from in the</u> library <u>provided in (i)</u> with an oligonucleotide, to form a plurality of peptoid-oligonucleotide mixtures, wherein said oligonucleotide is between about 10 and 50 nucleotides in length;
 - (iii) contacting each said mixture with a cell;
- (iv) screening each cell for transfection of the oligonucleotide, to identify transfected cells; and
 - (v) identifying transfecting peptoids in mixtures contacted with transfected cells.
- 14. (Original) The method of claim 13, wherein said library of peptoids is provided in an array of physically separated compartments.
- 15. (Previously presented) The method of claim 14, wherein said peptoids are supported on solid particles in said physically separated compartments.
- 16. (Previously presented) The method of claim 15, further comprising the step of releasing the peptoids from the particles in said compartments, prior to said contacting step (ii).
- 17. (Original) The method of claim 15, wherein each compartment contains a single particle, and each particle contains a single peptoid.

18-20 (Canceled)

21. (Previously presented) The method of claim 13, wherein, in step (iii), each said mixture is contacted with a plurality of distinct cell types.

22-24. (Canceled)

25. (Previously presented) The method of claim 13, wherein in formula I, R^a comprises a lipid moiety, and R^c is selected from -NH₂, -NHR, and -NH(C=O)R, where R is lower alkyl.

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- 26. (Original) The method of claim 25, wherein said lipid moiety is a sterol.
- 27. (Previously presented) The method of claim 13, wherein in formula I, each of R¹ and R² is hydrogen.
- 28. (Previously presented) The method of claim 13, wherein in formula I, at least one R^b includes a group which is cationic at physiologically relevant pH, and at least one R^b is uncharged at physiologically relevant pH.
- 29. (Previously presented) The method of claim 28, wherein said cationic group is selected from aminoalkyl, ammonium, guanidino, amidino, imidazole, and pyridinium.
 - 30-32 (Canceled)
- 33. (Currently amended) The method of claim 13, wherein said peptoids having a plurality of unknown different sequences have the general formula II:

$$R^{a} - (N(R^{b1}) - C(R^{1})(R^{2}) - C(O) - N(R^{b2}) - C(R^{1})(R^{2}) - C(O) - N(R^{b3}) - C(R^{1})(R^{2}) - C(O))_{h} - R^{0}$$

where

R^{b1} is a cationic moiety, R^{b2} is a non-cationic moiety, R^{b3} is a non-cationic moiety at physiologically relevant pH; and

n is an integer selected from 2 to about 16, and wherein

the sequences of individual peptoids in the provided library are unidentified prior to transfection screening.

- 34. (Previously presented) The method of claim 13, wherein said peptoids comprise at least two different cationic moieties.
- 35. (Currently amended) The method of claim 13, wherein <u>providing</u> said library <u>comprises</u> <u>synthesizing the library is synthesized</u> by a mix-and-split protocol.

- 36. (Previously presented) The method of claim 13, wherein identifying transfecting peptoids comprises determining their sequence.
- 37. (Currently amended) The method of claim 36, wherein the peptoid sequence is determined by tandem mass spectrometry a mass-spectrographic method.